

## CLAIM AMENDMENTS

1. (currently amended): A process for preparing an  $\alpha$ -aminonitrile with enhanced optical purity which process comprises  
contacting a mixture of the enantiomers of a chiral ~~N-formyl~~  $\alpha$ -aminonitrile with an acylase selective for one of the enantiomers,  
wherein said mixture is in the N-formyl form so that ~~whereby~~ one of the enantiomers of the said N-formyl- $\alpha$ -aminonitrile is selectively deformylated to obtain ~~into~~ the unprotected corresponding unprotected  $\alpha$ -aminonitrile; or  
wherein said mixture is in the unprotected form and a formylating agent is provided so that one of the enantiomers is selectively converted to the corresponding N-formyl  $\alpha$ -aminonitrile.
2. (canceled)
3. (currently amended): The process of claim 1 ~~claim 2~~ wherein the formylating agent is formic acid, a formic acid amide or a formic acid ester.
4. (previously presented): The process of claim 1, wherein the acylase is a peptide deformylase having a bivalent metal ion cofactor from group 5-11 of the periodic system.
5. (currently amended): The process of claim 1, wherein the peptide deformylase is of ~~chosen from~~ the class EC 3.5.2.27 or EC 3.5.1.31.
6. (previously presented): The process of claim 1, wherein the peptide deformylase contains the sequences (I) HEXXH, (ii) EGCLS and (iii) GXGXAAXQ.
7. (previously presented): The process of claim 4, wherein the peptide deformylase is from *Escherichia coli*.
8. (previously presented): The process of claim 4, wherein the bivalent metal is Fe, Ni, Mn or Co.

9. (previously presented): The process of claim 8, wherein the bivalent metal is Ni.

10. (previously presented): The process of claim 1, which further comprises adding a stabilisation agent.

11. (previously presented): The process of claim 10 wherein the stabilisation agent is catalase.

12. (previously presented): The process of claim 10 wherein the bivalent metal is Fe.

13-21 (canceled)